



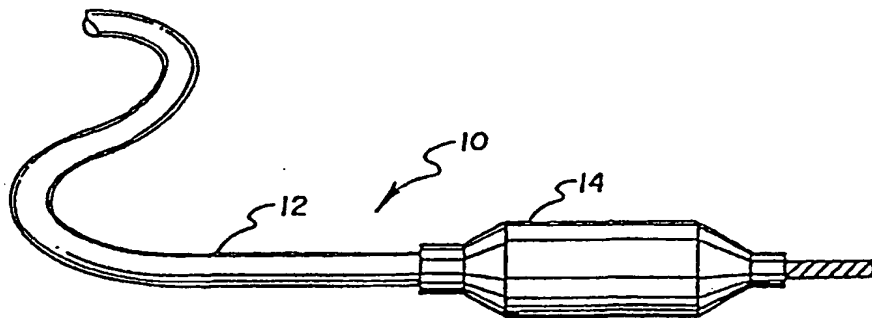
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : A61L 29/00	A2	(11) International Publication Number: WO 00/50105 (43) International Publication Date: 31 August 2000 (31.08.00)
(21) International Application Number: PCT/US00/03821 (22) International Filing Date: 15 February 2000 (15.02.00) (30) Priority Data: 09/257,677 25 February 1999 (25.02.99) US (71) Applicant: SCIMED LIFE SYSTEMS, INC. [US/US]; One Scimed Place, Maple Grove, MN 55311-1566 (US). (72) Inventors: WANG, Lixiao; 12822 86th Place North, Maple Grove, MN 55369 (US). CHEN, Jianhua; 4725 Terraceview Lane, Plymouth, MN 55446 (US). (74) Agent: STEINKRAUS, Walter; Vidas, Arrett & Steinkraus, 6109 Blue Circle Drive, Suite 2000, Minnetonka, MN 55343-9131 (US).		(81) Designated States: CA, JP, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>Without international search report and to be republished upon receipt of that report.</i>

(54) Title: MEDICAL DEVICES MADE FROM POLYMER BLENDS CONTAINING LOW MELTING TEMPERATURE LIQUID CRYSTAL POLYMERS

(57) Abstract

A medical device, at least a portion of which is composed of a polymeric material in which the polymeric material is a melt blend product of at least two different thermoplastic polymers, one of the thermoplastic polymers being a thermoplastic liquid crystal polymer (LCP) having a melting point of less than 250 °C. The portion of the device made from the melt blend may be a catheter body segment or a balloon for a catheter. The LCP blends suitably also include a non-LCP base polymer having a melting point in the range of about 140 °C to about 265 °C.



FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakistan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

MEDICAL DEVICES MADE FROM POLYMER BLENDS CONTAINING LOW MELTING TEMPERATURE LIQUID CRYSTAL POLYMERS

U.S. Application No. 09/257,677 from which this application claims
5 priority, is incorporated in its entirety herein by reference.

BACKGROUND OF THE INVENTION

In copending US application 08/926,905 (corresponding to
PCT/US98/18345 filed Sept. 4, 1998) there are described medical balloons made from
10 liquid crystal polymer blends. The blends comprise polymer melt blend product of

- a) a thermotropic main-chain liquid crystal polymer (LCP);
- b) a crystallizable thermoplastic polymer; and
- c) at least one compatibilizer for a) and b).

The melt blend balloons so produced have very high strength, but have relatively low
15 compliance and flexibility.

The practice of the invention of application 08/926,905, however, has
been limited in that the thermoplastic polymer was a material with a relatively high
melting temperature, such as crystallizable polyester or polyamide polymers. The
known LCPs had melting points above 275°C, thus requiring that the thermoplastic
20 polymer be stable at temperatures near or above the LCP melting temperature in order to
process the melt blend.

Many thermoplastic polymers have higher flexibility and elasticity than
polyesters or polyamides but their melting points have been too low to be processable in
melt blends with LCPs.

25 Recently LCPs with melting points below 250°C have been prepared and
commercialized. The inventors of the present invention have now discovered a much
wider range of thermoplastic polymers can be blended with such low melting
temperature LCPs to produce blend materials useful in fabricating medical devices.

SUMMARY OF THE INVENTION

In one aspect the invention comprises a medical device at least a portion of which is composed of a polymeric material in which the polymeric material is a melt blend product of at least two different thermoplastic polymers, one of the thermoplastic polymers being a thermoplastic liquid crystal polymer having a melting point of about 275°C or less, and especially 250°C or less. Catheters and catheter balloons are specific medical devices to which the invention may be applied.

The low temperature LCP component may be used at relatively low levels to impart higher strength and resistance to shrinkage to base polymer materials of greater flexibility, softness or elasticity than had previously been usable with available LCPs.

DESCRIPTION OF THE DRAWING

Fig. 1 is a perspective fragmentary view of a balloon catheter embodiment of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

The blend products used in the present invention include a thermoplastic non-LCP base polymer in an amount of from about 50 to about 99.9% by weight, preferably from about 85 to about 99.5 percent. The blend products also include from about 0.1 to about 20 weight percent, more preferably from about 0.5 to about 15 percent, of a liquid crystal polymer having a melting point of less than 275°C, preferably less than 250°C. A melt compatibilizer, such as disclosed in application 08/926,905, may also be employed in an amount of from 0 to about 30 weight percent.

The base polymer should have a melting point within about 70°C, preferably within about 50°C and more preferably within about 35°C of the liquid crystal polymer component. Suitably the base polymer has a melting point in the range of from about 140°C to about 265°C, preferably about 220°C or less, and more preferably from about 150°C to about 210°C. Depending on the liquid crystal polymer melting temperature, the base polymer may be for instance an acetal homopolymer or copolymer (typical mp 160-185°C); cellulosic polymers (mp. 140-190°C); poly(chlorotrifluoroethylene) (mp. 200-220); poly(vinylidene fluoride) (mp 155-180°C);

nylon 6,6 (mp. 250-260); nylon 6 (mp 215-225); nylon 6,10 (mp 210-220); nylon 12 (mp 170-180); nylon 11 (mp 180-190); polyoxymethylene (mp 165-185); higher melting grades of poly(methyl methacrylate) (e.g. mp 140-160°C); polypropylene homopolymers and copolymers (mp 160-175); polycarbonate polymers and copolymers (mp 220-5 230°C); poly(ethylene-vinyl alcohol) (mp 140-180); polyethylene terephthalate; polybutylene terephthalate; polytrimethylene terephthalate; thermoplastic polyurethanes (aromatic and/or aliphatic); thermoplastic elastomers such as polyester elastomers sold under the tradenames Hytrel® and Arnitel®, polyamide elastomers sold under the tradename Pebax®, and thermoplastic polyurethane elastomers sold under the tradename 10 Pellethane®. Particularly preferred base polymer materials include Pebax® 7033 (mp 174°C) and 7233 (mp 175°C), sold by Atochem North America, and Arnitel EM 740 (mp 221°C), sold by DSM Engineering Plastics.

Use of some of these base polymers in LCP blends has been described in the prior application 08/926,905, for instance PET/LCP blends. However, by using 15 lower melting temperature LCPs, as described herein, processing is made easier. For instance, where there is a large temperature difference between the base polymer and the LCP component, a dual extruder may have had to be used to allow the polymers to be separately melted before they could be mixed. With a smaller difference in melt temperatures the melt blend of LCP and base polymer can be prepared by melting a dry 20 blend of the two polymers, or one of the two polymers in solid form may be added to a melt of the other, without substantial polymer degradation. A dual extruder technique can still be used to obtain blends with base polymers whose melt temperature is substantially lower than that of the LCP used in the present invention. Therefore the range of usable base polymers is substantially increased in the present invention over those of prior 25 application 08/926,905.

The LCP used in the invention hereof is one characterized by a melting point below 275°C, preferably below 250°C, suitably in the range of 150-249°C, and even more preferably about 230°C or less. The LCP is suitably a thermotropic liquid crystal polymer. Specific such LCPs include Vectra® LKX 1107, a polyester-type liquid 30 crystal polymer (mp 220°C), and Vectra® LKX 1111, a polyesteramide-type liquid crystal polymer (mp. 220°C), both sold by Ticona, a Hoechst company.

Compatibilizers also may be used in the melt blend composition. The compatibilizer may be for instance a block copolymer comprising a block which is structurally similar or otherwise is soluble in the base polymer and a block which is structurally similar or otherwise soluble with the LCP. Compatibilizers may be
5 necessary if phase separation of the blend in the melt phase is a problem. However, phase separation of the solid phase melt blend product is not necessarily a reason to employ a compatibilizer. Solid phase separation may enhance the reinforcing effect of the LCP component. Optical clarity, however, is lost with phase separation in the solid phase. Use of a compatibilizer may be useful if optical clarity is a desired objective or
10 where it is desired to improve adhesion between LCP fiber and the base polymer.

The blend materials described herein are particularly suited for use in forming dilatation and/or stent placement catheters or balloons thereon. Such catheters are used for percutaneous transluminal angioplasty and other minimally invasive procedures. Use in forming a proximal or intermediate portion of the catheter body may
15 reduce or eliminate the need for braid or other physical reinforcement so that a reduced profile may be provided.

A particularly preferred use of the melt blend materials described herein is as a material for a catheter balloon. The balloon diameter may be from about 1.5 to about 30 mm, depending on the application to which it is put, and are suitably formed to
20 provide a double wall thickness, measured on the uninflated collapsed balloon, of about 0.0002" - 0.0020".

The balloons of the invention may be either single layer balloons, or multilayer balloons.

25 Referring to the drawing, there is shown in Figure 1 a catheter 10 comprising an elongated flexible tube 12 with a balloon 14, made of an LCP reinforced polymer blend in accordance with the invention hereof, mounted at the distal end thereof. A portion of tube 12 also may be formed from an LCP reinforced polymer blend, which may be the same or different from the blend used to form the balloon.

30

Balloon formation may be begun by extruding a tube from a melt of the

polymer blend material. Some initial orientation of the LCP occurs as the blend material is drawn down during the extrusion process. This process is typically known as machine orientation and is in the direction of the extrusion operation. Orientation which occurs during the extrusion process is desirable as it induces formation of fiber form LCP in the tubing so-formed. Orientation can be enhanced by increasing extrudate puller speed. Also, if an angled fiber morphology is desired, a counter-rotating die and mandrel system can be used in the extrusion.

Following extrusion, the extruded tube optionally may be conditioned at 20-30°C at a controlled humidity in the range of 10-50% for a period of at least 24 hours. This conditioning provides a constant low moisture level in the tube which prevents hydrolysis and helps to optimize the orientation of the polymer in the subsequent blowing steps.

Balloon blowing may follow conventional single or multi-step techniques known in the art, for instance free blowing, mold blowing, or a combination of both, optionally with a preceding axial stretching step. The axial stretch ratio, if used, is suitably from about 2x to about 5x. Balloon forming will typically be performed at a temperature in the range of 95°C to 165°C, depending on the base polymer material and the amount of LCP incorporated into the blend. The balloon forming step should be performed above the glass transition temperature but below the melt temperature of the base polymer material (for block copolymers the blowing temperature should be above the highest glass transition). The radial expansion ratio is suitably from about 3x to about 12x. Depending on the technique, expansion pressures may range from about 200-500 psi (1379 -3447 kPa).

In some cases it may be desirable to subject the formed balloon to a heat set step. In this step the pressurized balloon is held for a brief time, suitably about 5-60 seconds, at a temperature above that used to form the balloon after which the mold is rapidly quenched to ambient temperature and the balloon removed from the mold.

In the absence of a compatibilizer, or where the compatibilizer is only effective to compatibilize the melt, the LC and base polymers will typically undergo phase separation on cooling so that an opaque article is obtained. The phase separation, however, occurs on a microscopic scale so that the LC discontinuous phase is uniformly

distributed in a continuous base polymer phase. The LC discontinuous phase is fibrous, and the fibers orient during the stretching and blowing steps of the balloon formation so a high level of reinforcement is provided to the base polymer. However, reinforcement by the fibrous LC phase can be achieved without a major reduction in flexibility and without presenting huge increases in melt viscosity, both of which effects are commonly encountered when reinforcing fillers are added to thermoplastic polymer compositions. Moreover, the fiber size is so small that, even with the extremely thin films encountered in angioplasty balloons, film porosity is not created.

The invention is illustrated by the following non-limiting examples.

EXAMPLES

EXAMPLE 1

Pebax 7033 polymer was melt blended at a temperature of 225°C with liquid crystal polymer Vectra LKX 1107 at the ratio of 95% to 5% respectively by weight and the mixture was extruded into tubing of 0.018 x 0.037 inch (0.48 x 0.94 mm). A 3.0 mm balloon was formed from the tube at 98°C and at 450 psi (4102 kPa) forming pressure using a 3.0 mm mold form in a single blowing step. The balloon had a double wall thickness of 0.00175 inch (0.044 mm) and had an opaque appearance. The balloon burst at 265 psi (1827 kPa). This reinforced composite balloon has much higher puncture resistance and more durability than a similar balloon made from 100% Pebax 7033.

Improved length stability upon expansion is a desirable property for high strength, relatively compliant balloons used for stent deployment. The following Examples 2 and 3 demonstrate that the LCP blends used in the invention provide improvement in length stability for such balloons.

EXAMPLE 2

The same composition as shown in Example 1 was used to extrude a tube of 0.022 x 0.036 inch (0.56 x 0.91 mm). The 3.0 mm balloon was made at 95°C with a blowing pressure of 400 psi (2758 kPa). The balloon with double wall thickness of

0.0014 inch (0.036 mm) was inflated from 4 atm (405 kPa) to 13 atm (1317 kPa) at 1 atm (101 kPa) increments and the balloon length change was 2.5% at the span of 4-13 atm.

For comparison 100% Pebax 7033 tubing with dimension of 0.0192 x 0.0344 (0.49-0.87 mm) was used to form 3.0 mm balloon at 95°C and 400 psi (2758 kPa) blowing pressure. The formed balloon with double wall thickness of 0.0014 inch (0.036 mm) was inflated from 4 atm (405 kPa) to 13 atm (1317 kPa) at 1 atm (101 kPa) increments and the balloon grew 8.0% of its original length before inflation.

10 EXAMPLE 3

The same molding conditions as in the previous examples were used for this example. A 40 mm long 3.0 mm diameter balloon mold was used to make a 100% Pebax 7033 balloon. The formed balloon had a body length of 37.0 mm after the balloon was removed from the mold. The same mold and balloon forming conditions were used for a LCP reinforced Pebax 7033 balloon formed from the melt blend product described in Example 1. The formed balloon had the body length of 38.5 mm, corresponding to a 50% improvement in balloon body length stability as a result of the inclusion of the 5% LCP component.

The foregoing examples and disclosure are intended to be illustrative and not exhaustive. These examples and description will suggest many variations and alternatives to one of ordinary skill in this art. All these alternatives and variations are intended to be included within the scope of the attached claims. Those familiar with the art may recognize other equivalents to the specific embodiments described herein which equivalents are also intended to be encompassed by the claims attached hereto.

CLAIMS

What is claimed is:

1. A medical device at least a portion of which is composed of a polymeric material in which the polymeric material is a melt blend product of at least two different thermoplastic polymers, one of the thermoplastic polymers being a thermoplastic liquid crystal polymer (LCP) having a melting point of less than 250°C.
2. A device as in claim 1 wherein the medical device is a catheter.
3. A device as in claim 2 wherein said device portion is a balloon mounted on the catheter.
4. A device as in claim 1 wherein the melt blend product includes said LCP in an amount of about 0.1 to about 20 weight percent and a thermoplastic non-LCP base polymer in an amount of from about 50 to about 99.9% by weight, the base polymer having a melting point in the range of about 140°C to about 265°C.
5. A device as in claim 4 wherein the base polymer has a melting point of about 220°C or less.
6. A device as in claim 5 wherein the melting point of the base polymer is from about 150° to about 210°C and the melting point of the LCP is about 150°C to about 230°C.
7. A device as in claim 4 wherein the base polymer is selected from the group consisting of acetal homopolymers and copolymers, cellulosic polymers, poly(chlorotrifluoroethylene), poly(vinylidene fluoride), nylon 6,6, nylon 6, nylon 6,10, nylon 12, nylon 11, polyoxymethylene, poly(methyl methacrylate) having a melting point in the range of above 140°C, polypropylene homopolymers and copolymers, polycarbonate polymers and copolymers, poly(ethylene-vinyl alcohol), polyethylene terephthalate, polybutylene terephthalate, polytrimethylene terephthalate, thermoplastic

polyurethanes (aromatic and/or aliphatic) and thermoplastic elastomers.

8. A device as in claim 7 wherein said base polymer is a thermoplastic polyamide elastomer or a thermoplastic polyester elastomer.

5

9. A device as in claim 8 wherein said base polymer is present in said melt blend in an amount of from about 85 to about 99.5 weight percent and said LCP is present in an amount of 0.5 to about 8 percent.

10 10. A balloon for a medical device, the balloon being prepared by radial expansion of a tubular parison of polymeric material, wherein the polymeric material is a melt blend product of at least two different thermoplastic polymers, one of the thermoplastic polymers being a thermoplastic liquid crystal polymer (LCP) having a melting point of less than 250°C

15

11. A balloon as in claim 10 wherein the melt blend product includes said LCP in an amount of about 0.1 to about 20 weight percent and a thermoplastic non-LCP base polymer in an amount of from about 50 to about 99.9% by weight, the base polymer having a melting point in the range of about 140°C to about 265°C.

20

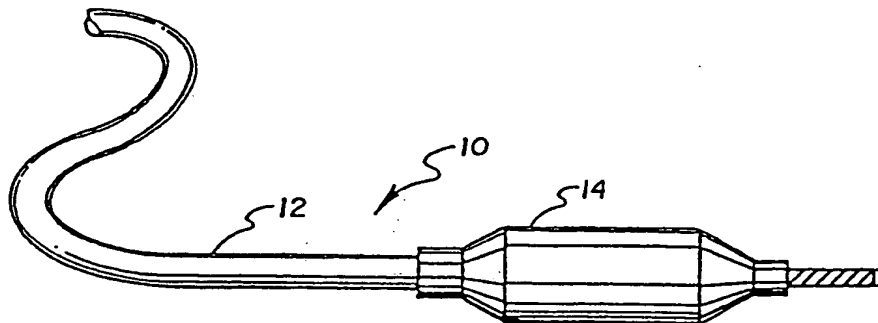
12. A medical device, at least a portion of which is composed of a polymeric material, in which the polymeric material comprises at least two different thermoplastic polymers, one of the thermoplastic polymers being a thermoplastic liquid crystal polymer (LCP) and a second of the thermoplastic polymers being a non-LCP base polymer, the
25 polymeric material being a two-phase system of LCP fibers distributed in the non-LCP base polymer.

13. A medical device as in claim 12 wherein said medical device portion is an elongated structure, and the fibers are oriented in the longitudinal direction of the
30 structure.

14. A medical device as in claim 12 wherein said medical device portion is an elongated structure, and the fibers are oriented at an angle to the longitudinal direction of the structure.
- 5 15. A medical device as in claim 12 wherein the LCP has a melting point of less than 275°C and the base polymer has a melting point in the range of about 140°C to about 265°C.
16. A medical device as in claim 12 wherein the base polymer is a
10 thermoplastic elastomer.
17. A medical device as in claim 12 wherein said medical device portion is a catheter balloon.
- 15 18. A method of forming a balloon by radial expansion of an extruded tubular parison of a polymer material comprising a thermoplastic non-LCP base polymer, the method comprising:
- melt blending said non-LCP base polymer with 0.1 to 20 weight % of an LCP prior to formation of said parison;
- 20 extruding the parison in a manner so that the LCP phase separates during solidification of the melt blend product and forms longitudinally oriented fibers in a matrix of said base polymer; and then
- radially expanding the parison to form said balloon.
- 25 19. A method as in claim 18 wherein the LCP has a melting point of less than 275°C and the base polymer has a melting point in the range of about 140°C to about 265°C.
20. A method as in claim 19 wherein the LCP melting point is in the range of
30 150°C to 249°C.

1/1

Fig.1



THIS PAGE BLANK (USPTO)

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
31 August 2000 (31.08.2000)

PCT

(10) International Publication Number
WO 00/50105 A3

(51) International Patent Classification⁷: A61L 29/06,
29/14

(74) Agent: STEINKRAUS, Walter; Vidas, Arrett &
Steinkraus, 6109 Blue Circle Drive, Suite 2000, Min-
netonka, MN 55343-9131 (US).

(21) International Application Number: PCT/US00/03821

(81) Designated States (*national*): CA, JP.

(22) International Filing Date: 15 February 2000 (15.02.2000)

(25) Filing Language: English

(84) Designated States (*regional*): European patent (AT, BE,
CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,
NL, PT, SE).

(26) Publication Language: English

(30) Priority Data:
09/257,677 25 February 1999 (25.02.1999) US

Published:
— With international search report.

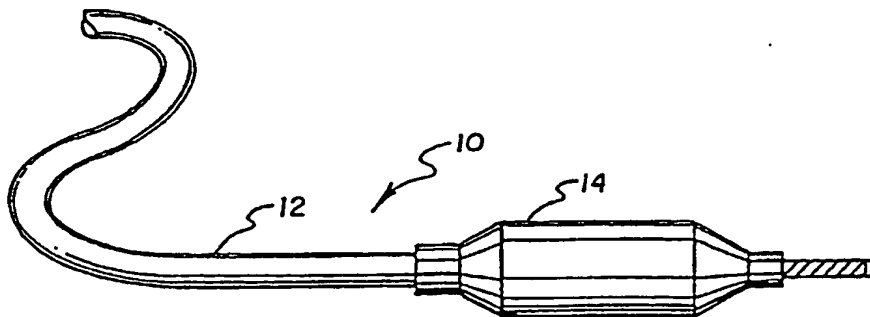
(71) Applicant: SCIMED LIFE SYSTEMS, INC. [US/US];
One Scimed Place, Maple Grove, MN 55311-1566 (US).

(88) Date of publication of the international search report:
14 December 2000

(72) Inventors: WANG, Lixiao; 12822 86th Place North,
Maple Grove, MN 55369 (US). CHEN, Jianhua; 4725
Terraceview Lane, Plymouth, MN 55446 (US).

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: MEDICAL DEVICES MADE FROM POLYMER BLENDS CONTAINING LIQUID CRYSTAL POLYMERS



(57) Abstract: A medical device, at least a portion of which is composed of a polymeric material in which the polymeric material is a melt blend product of at least two different thermoplastic polymers, one of the thermoplastic polymers being a thermoplastic liquid crystal polymer (LCP) having a melting point of less than 250 °C. The portion of the device made from the melt blend may be a catheter body segment or a balloon for a catheter. The LCP blends suitably also include a non-LCP base polymer having a melting point in the range of about 140 °C to about 265 °C.

WO 00/50105 A3

INTERNATIONAL SEARCH REPORT

International Application No

PC1, JS 00/03821

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61L29/06 A61L29/14

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61L A61M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 92 08512 A (BOSTON SCIENT CORP) 29 May 1992 (1992-05-29) page 5, line 24 - line 30 claims 1,8,9,13,14,25 ---	1-11
X	EP 0 448 886 A (BECTON DICKINSON CO) 2 October 1991 (1991-10-02) page 4, column 25 -column 40 -----	1-11

☐ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

* Special categories of cited documents :

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

& document member of the same patent family

Date of the actual completion of the international search

16 June 2000

Date of mailing of the international search report

19. 09. 2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Heck, G

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US 00/03821

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-11

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

1. Claims: 1-11

Medical devices at least partially made of a polymer blend of a liquid crystal polymer with a melting point of less than 250AC and at least one more thermoplastic polymer.

2. Claims: 12-20

Medical devices at least partially made of a polymer blend of a liquid crystal polymer and at least one more non-liquid crystal thermoplastic polymer in the form of fibers distributed in the non-liquid crystal polymer.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCi, JS 00/03821

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9208512 A	29-05-1992	CA 2094597 A	10-05-1992
		DE 69131104 D	12-05-1999
		DE 69131104 T	14-10-1999
		EP 0556309 A	25-08-1993
		EP 0974370 A	26-01-2000
		JP 6504458 T	26-05-1994
		US 5306246 A	26-04-1994
EP 0448886 A	02-10-1991	AT 123231 T	15-06-1995
		AU 631949 B	10-12-1992
		AU 6792390 A	03-10-1991
		CA 2031805 A	27-09-1991
		DE 69019816 D	06-07-1995
		DE 69019816 T	11-01-1996
		ES 2075175 T	01-10-1995
		IE 67657 B	17-04-1996
		JP 2096874 C	02-10-1996
		JP 4221571 A	12-08-1992
		JP 8011129 B	07-02-1996
		NZ 236404 A	23-12-1992
		US 5453099 A	26-09-1995
		US 5226899 A	13-07-1993

THIS PAGE BLANK (USPTO)